## REMARKS

Entry of the claim and specification amendments are requested because it is believed that such amendment put the application in condition for allowance. It is also believed that a new search would not be required and that only minimal reconsideration would be necessary. Entry of the amendments is also believed to place the application in better condition for appeal.

Claims 28, 39, 51 and 52 have been rejected under 35 U.S.C. 102(b) as being anticipated by Heyes et al. (U.S. Patent No. 5,093,208). Applicants traverse this rejection.

Applicants coextrude the polyolefin (i.e., polypropylene or polyethylene) and the adhesion-promotion agent. The melted materials exit the coextruder nozzle. The coextrudate is still very hot for a substantial time after exiting the nozzle. These are inherent features of the coextrusion.

When the very hot coextrudate is combined with the aluminum foil, the temperature of the aluminum has to be low enough so that the outer surface of the polyolefinic portion of the coextrudate stays below the crystallite melt point  $(T_K)$  of the olefin. The coextrudate is combined with the aluminum foil between two rollers. The aluminum foil acts as a heat sink re the coextrudate – this is inherent – thereby maintaining the temperature of the outer surface of the plastic coating below the crystallite melt point  $(T_K)$  of the olefinic plastic.

Nowhere does Heyes et al. teach or suggest using the aluminum foil as a heat sink. Heyes et al. teaches preheating the aluminum foil to above or near the melting point of the polyester - that then results in raising (not lowering) the

temperature of the cooler polyester coating. Figures 1 and 2 show the polyester in rolls at room temperature. Heyes et al. uses the heated aluminum as a heat source. Heyes et al. does not anticipate any of applicants' process claims.

The invention of Heyes et el. is limited to using extrudates or coextrudates of polyester or polyesters. On this ground alone Heyes et al. does not anticiapte any of applicants' process claims.

Heyes et al. discloses that polyolefin coatings on metal sheet (aluminum foil) are unsatisfactory for the manufacture of drawn and wall-ironed cans (DWI cans). Examples 11 and 12 of Heyes et al. are comparative examples of prior art to the Heyes et al. invention. Examples 11 and 12 use laminate types H and I, respectively, that each has coextruded polypropylene composite film on both sides of the aluminum sheet. (The laminates of both Examples 11 and 12 are stated to exhibit poor formality and to give metal failure in can forming.)

Examples 11 and 12 of Heyes et al. use the Heyes et al. method of laminate formation that includes using "preheating the metal strips" and then passing the coextruded polyolefin composite films and the preheated metal strips into a pair of nip rolls. Accordingly, Examples 11 and 12 of Heyes et al. do not anticipate any of the applicants' process claims.

No where does Heyes et al. teach the use of aluminum sheet that has a temperature lower than the coextruded polyester film. Heyes et al. generically discloses laminating the coextruded polyester film to the metal sheet but such disclosure is not anticipatory because it does not teach all of the requirements and limitations of applicants' process claims. Heyes et al.'s generic disclosure is

not a teaching of a process step where the relatively cool temperature of the aluminum foil causes the outer surface of the hot polyolefin to stay below its crystaline melt point ( $T_K$ ). Heyes et al. states:

"The laminated metal sheet of the invention is prepared by a process which compromises adhering directly to one or both major surfaces of the metal sheet a film comprising a polyester, the lamination conditions being such that during lamination the polyester film or films in the metal/polymer laminate is or are converted into non-crystalline or amorphous form." [Col. 3, lines 33 to 39]

The only specific schemes of Heyes et al. are ones which use preheating of the aluminum sheet. The preheating is not identified by Heyes et al. as being the preferred feature of the two preferred process – each of the two preferred processes of Heyes et al. involve all of such process, not part thereof.

Heyes et al. states:

"In one preferred process of preparing the metal polymer laminates in accordance with the invention polyester monolayer film or films are adhered to the metal sheet by heating the metal sheet to a temperature (T1) above the melting point of the polyester films, the temperature (T1) being such that during lamination of the polyester films to the metal sheet, the outer surfaces of the polyester films remain below their melting points, \*\*\*."

"In an alternative preferred process, the polyester film or films are composite films (A) comprising an inner layer (A1) and an outer layer (A2),

and the composite polyester films are simultaneously adhered to the metal sheet by a process which comprises

(1) heating the metal sheet to a temperature (T1) above the softening point of the polyester inner layer (A1) but below the melting point of the outer layer (A2), \*\*\*," [Emphasis supplied] [Col. 3, lines 40 to 62]

In the disclosure of Heyes et al. that deals specifically with the temperature of the coextruded plastic coatings, the coextruded plastic materials are at a temperature less than the preheated metal (aluminum) sheet. Heyes et al. states:

"Polymer/metal/polymer laminates were prepared by a lamination process performed in apparatus as illustrated schematically in FIG. 1 or FIG. 2 of the accompanying drawings. A metal sheet M was pre-heated by infrared or induction heating to an appropriate temperature T<sub>1</sub> by a heater 1. Temperature T<sub>1</sub> is usually within the range 140° and 350°C.

Polyester films A and B were fed from feed rolls 2 and 4 and laminated to the opposite sides of the pre-heated metal sheet between lamination rolls 6, 8, \*\*\*." [Emphasis supplied] [Col. 6, line 65, to Col. 7, line 5]

Figures 1 and 2 of Heyes et al. do not show any preheating of polyester films A and B, either rolled up or being fed to lamination rolls 6, 8. Since Heyes et al. does not recite any temperature for polyester films A and B before the lamination steps (rollers 6, 8), in accordance with scientific/technical practice, polyester films A and B were at room temperature.

The Office Action stated that, as to Claims 28 to 52, Heyes et al. discloses a process for the production of an aluminum foil (see column 2, lines 15 to 24) coated with a sealable and sterilizable plastic based on polypropylene (PP) comprising co-extruding the plastic maleic anhydride (MAH) graft modified PP (an adhesion promoting agent) and combining co-extruded PP composite with the aluminum foil between two rollers (see Fig. 1; Table I, type H; column 8, lines 9 to 19), the temperature at the outer surface of the plastic lies below the (crystallite) melt point  $(T_K)$  of the plastic (see column 3, lines 25, 46 and 47), then passing continuously the coated aluminum foil through a heater 10 (oven) to increase the adhesion strength between the aluminum foil and the plastic coating (see Fig. 1; column 7, lines 12 to 16) with a temperature set so that the temperature at the outer surface of the plastic lies above the (crystallite) melt point (T<sub>K</sub>) of the plastic (see column 3, lines 48 to 51), and quenching (cooling in a shock-like manner) the coated aluminum foil such that the crystalline plastic is converted to non-crystalline or amorphous form (i.e., crystalline proportion at least in the surface area of the cooled PP layer is as small as possible) (see column 1, lines 45 to 47; column 2, lines 1 to 15; column 3, lines 39). Applicants traverse this statement as being an incorrect statement of the disclosure of Heyes et al. Specifically, Heyes et al. only discloses generically adhering a polyester to a metal sheet, without any reference to the temperature of either, or specifically adhering a polyester (or polyolefin) film to a metal sheet that has been preheated, with specific disclosure showing that the polyester film is at

room temperature (that is below the preheating temperature of the metal sheet).

Heyes et al. is not an anticipatory reference.

The Office Action stated that it is the Examiner's position that the surface area of the cooled PP layer has claimed properties such that if the quenched non-crystalline plastic still has small amounts of crystals, then the crystal grains are as small as possible *inherently* since it is produced by a method identical or substantially identical processes to that of claimed invention. Applicants traverse this statement. The process of Heyes et al. and the process of the applicants' claims are not identical or substantially identified, as shown above. The difference between preheating the aluminum sheet and the coextruded polyolefin/adhesion-promotion agent results in substantial differences. The Examiner's assertion of inherency is faulty and lacks factual support.

Heyes et al. states:

"Examples 11 and 12 show that laminates formed from polypropylene materials of the type described in GB 2003415 exhibited poor formality. Such laminates were found to give metal failure can forming." [Emphasis supplied] [Col. 9, lines 65 to 68]

Table II also recites poor formability for Examples 11 and 12.

So it is clear that Heyes et al. does not inherently achieve crystal grains as small as possible.

The feature of applicants' process of using hot coextruded polyolefin/adhesion-promotion agent and cooler aluminum foil helps provide different results. The hot/soft polyolefin/adhesion-promotion agent of applicants'

process has more time in the nip region to effect elevated temperature adhesion to the aluminum foil than does the scheme of Heyes et al. wherein the plastic composite has to first be heated up by the preheated aluminum sheet during the very short time period involved in passing through the two laminated rollers. The Examiner's attempt to use the concept of inherency fails.

Applicants' process produces containers that have essentially no white breaks in the deformation area.

The Office Action stated: that it is held that where the claimed and prior art products are identical or substantially identical in structure or composition, or are produced by identical or substantially identical processes, claimed properties or functions are presumed to be inherent; (see MPEP 2111.02 and 2112.01, and In re Best 562 F.2d 1252, 1255, 195 USPQ 430, 433 (CCPA 1977); and "When the PTO shows a sound basis for believing that the products of the applicant and the prior art are the same, the applicant has the burden of showing that they are not," In re Spada, 911 F.2d 705, 709, 15 USPQ2d 1655, 1658 (Fed. Cir. 1990). Applicants traverse this statement as being an incorrect statement of the pertinent decisions and legal principles. The Spada decision involves products, that are not claimed by applicants. Instead, applicants claim a one-step production process that substantially differs from the multi-step production process of Heyes et al. There are actual physical differences in the processes that are not ones "claimed in terms of function property or characteristics." As shown above, the products of the two processes differ. The burden of proof has not shifted to applicants.

The Examiner has not factually established in the record that a prima facie showing of anticipation exists. The two processes are substantially different.

This rejection should be withdrawn.

The Office Action stated: that, as to Claim 29, the reheating temperature is  $270^{\circ}\text{C}$  (see Table II, type H; column 8, lines 9 to 19), which is at least  $20^{\circ}\text{C}$  above the crystallite melt point ( $T_K$ ) of PP since  $T_K$  of PP is around  $160^{\circ}\text{C}$ , as evidenced by applicants (see specification, page 3, line 33); and that, consequently, the temperature of heater 10 (oven) is also at least  $20^{\circ}\text{C}$  above the  $T_K$  of PP. Independent Claim 28 is not anticipated by Heyes et al., as factually shown above, so dependent Claim 29 is not anticipated by Heyes et al.

The Office Action stated that, as to Claim 51, the quenching (shock-like cooling) is carried out by <u>cold water</u> (see column 8, lines 6 to 8 and 19).

Independent Claim 28 is not anticipated by Heyes et al., as factually shown above, so dependent Claim 51 is not anticipated by Heyes et al.

This rejection should be withdrawn.

Not only is applicants' claimed process not anticipated by Heyes et al., it is unobvious over Heyes et al. The Examiner, who has the burden of proof, has factually established neither anticipation nor obviousness.

Applicants' process includes the step of coextruding the plastic (PP or PE) and the adhesion-promotion agent. The oleofinic plastic and the adhesion-promotion agent are melted in the extruder barrel. The New Encyclopedia Britannica, Macropaedia Volume 14, (1947), states:

"Extrusion. A major technique of the plastics industry, extrusion consists essentially of the melting and compression of plastic granules by the rotation of a screw conveyor in a long barrel to which heat and cold can be applied. The screw drives the plastic through a nozzle \*\*\*. \*\*\* so as to compress and generally homogenize the melting plastic. \*\*\*; it serves to complete the melting \*\*\* and \*\*\* pumping the molten plastic through the shaping nozzle." [Emphasis supplied] [Page 519]

Upon exiting from the coextruder, applicants' process moves the coextrudate rapidly to the two-rollers, were it is combined with the aluminum foil as both begin to pass through the two rollers. The aluminum foil acts as a heat sink.

One of the very core purposes of Heyes et al. is to provide an invention that does not use polyolefin coatings, that avoids the problems caused by the use of polyolefin coatings, and that provides advantages over polyolefin coatings. Heyes et al. states:

"It is known to use steel or aluminum coated with polyolefin coatings as a stock preparing DWI cans. Such materials are described, for example, in U.S. Pat. No 4,096,815 and British Patent 2003415; as far as we are aware, such materials have not found commercial application."

"We have found that polyolefin coatings do not form as well as thermoplastic polyesters." [Emphasis supplied] [Column 1, lines 14 to 21]

"Such [substantially non-crystalline or amorphous thermoplastic polyester] coatings out-perform polyolefin coatings in DWI can forming,

and retain better continuity and protection." [Emphasis supplied] [Column 1, lines 41 to 43]

Heyes et al. directs away from the use of polyolefin coatings and, hence, also directs away from applicants' claimed process.

As shown above, the prior art comparison Examples 11 and 12 of Heyes et al. (that combined polypropylene composite films and preheated aluminum sheet) provided "poor" formability and gave metal failure in can forming. One ordinarily skilled in the art is pointed away by Heyes et al. from the use of polypropylene coatings.

In the amendment (of August 28, 1991) in U.S.S.N. 07/642,566, upon which Heyes et al. issued, applicants submitted a copy of Koga et al., U.S. Patent No. 4,849,293, (and a copy of corresponding European Published European Patent Application 0262929) that had been cited in Heyes et al.'s corresponding U.K. application (the Examiner initialed references on Form PTO-FB-A820). The amendment, regarding Koga et al., states:

"Two particular amorphous polyester compositions are described.

A first composition comprises (A) a low crystalline modified polyolefin, (B) an amorphous polyester and (C) a silane coupling agent. A second amorphous polyester composition comprises (A), (B) and (C) together with (D) an inorganic filler. In both compositions (A) forms a 'matrix phase', and (B) forms 'a domain phase.' Composite laminates for damping materials are described (see Col. 6, lines 59 et seq. of '293). These include metal/polyester composite structures."

"There appears to be no disclosure of a process for making a laminate of metal and non-crystalline polyester which includes a step of providing a sheet of metal and a film of biaxially-oriented polyester having a semi-crystalline structure. Furthermore, there does not appear to be two separate heating steps followed by a rapid quenching step to form a laminate of metal and non-crystalline polyester. Additionally, there appears it be no disclosure of a composite polyester film comprising an inner polyester layer and an outer polyester layer."

"The matrix/domain phase composition of this document would be unlikely to be suitable in the manufacture of containers. If a polyolefin matrix phase were used in the manufacture of cans, it would not survive the drawing and wall ironing to which the laminates of the present invention are subjected." [Emphasis supplied] [Page 5, line 12, to page 6, line 3]

During the prosecution of Heyes et al., Heyes et al. further directed away from polyolefin coatings and away from applicants' claimed process.

Claims 30 to 50, 53 and 54 have been rejected under 35 U.S.C. 103(a) as being unpatenable over Heyes et al. (U.S. Patent No. 5,093,208) in view of Takano et al. (U.S. Patent No. 5,837,360). Applicants traverse this rejection.

Applicants have shown above that Heyes et al. does not make any of applicants' claims obvious. Takano et al. does not cure the defects of Heyes et al. in the search for applicants' claimed invention.

Takano et al. also directs one ordinarily skilled in the art away from applicants' claimed invention. In Takano et al., polypropylene and a modified polypropylene are melt-coextruded and laminated on at least one side of a preheated steel sheet, i.e., desirably preheated to a temperature of from 100°C to 160°C. Takano et al. asserts that any preheating below 100°C is unsatisfactory because then the laminated entity would be below the minimum temperature for initintion of the subsequent quenching. Note also that Takano et al. does not disclose laminating by passing the films and sheet through a pair of rollers.

Levendusky et al., i.e., U.S. Patent No. 5,919,517 (cited by the Examiner) in its background-of-the-invention section, states:

"U.S. Pat. No. 5,093,208 to Heyes et al. discloses a method for forming a laminated metal sheet in which a precast thermoplastic polyester film is pressed against one or both surfaces of a metal sheet to adhere the film to the sheet in a pressed against one or both surfaces of a metal sheet to adhere the film to the sheet in a non-crystalline form. The uncoated sheet of metal is heated to a temperature above the melting pot of the polyester film and the film is applied to the sheet under pressure to form a laminate material." [Emphasis supplied] [Column 1, lines 29 to 37]

(Levendusky et al. is assigned to ALCOA.) The art views Heyes et al. as preheating the uncoated aluminum sheet to a temperature above the melting point of the polyester film. The Examiner's attempt to stick Takano et al. into Heyes et al. would destroy the invention of Heyes et al.

The Offic Action stated that, as to Claims 30 to 35, 48 and 49, Heyes et al., as applied above, fails to teach that: T<sub>E</sub> of shock-like cooling is at least 40°C below T<sub>K</sub> of PP (Claims 30 and 48) or at least 60°C (Claim 31) or at least 80°C below T<sub>K</sub> of PP (Claim 32); the shock-like cooling speed (V<sub>A</sub>) is greater than 10°C/sec (Claims 33 and 49), greater than 50°C/sec (Claim 34) or greater than 100°C/sec (Claim 35). Heyes et al. does not teach or suggest any of applicants' claims, and Takano et al. does not cure the defects of Heyes et al. in the quest for applicants' invention.

The Office Action stated that Takano et al. teaches that the shock-like cooling of metal foil coated with co-extruded PP plastic/MAH modified PP (see column 2, lines 1, 54 to 55 and 61 to 67; column 3, lines 1 to 6) and heated above the T<sub>K</sub> of the PP based layers (see column 5, lines 35 to 52) with a speed (V<sub>A</sub>) greater than 10°C/sec (see column 6, lines 10 to 12), or greater than 50°C/sec, preferably greater than 100°C /sec (see column 6, lines 10 to 12), so that the T<sub>E</sub> of the shock-like cooling is at least 55°C (see column 5, lines 59 to 63), reduces crystallinity of PP based layers not more than 55 percent thereby providing the coated metal foil with superior corrosion reistance and peeling resistance (see column 1, lines 57 to 62; column 2, lines 9 to 11). This information does nothing to provide applicants' claimed invention from the attempted combination of the rejection references.

The Office Action stated that it would have been obvious to one of ordinary skill in the art at the time the invention was made to have carried out shock-like cooling of metal foil coated with co-extruded PP plastic/MAH modified

PP in Heyes et al. under conditions of Takano et al. with the expectation of providing the coated metal foil with desired superior corrosion resistance and peeling resistance as taught by Takano et al. Applicants traverse this statement. The combination of the two rejections does not result in applicants claimed invention. Also, both rejection references direct away from applicants' claimed invention.

The Office Action stated that, as to Claims 36, 39 to 43 and 50, Heyes et al. fails to teach that shock-like cooling is carried out by partial looping over at least one cooled roller (Claims 36 and 50) or by passing through ice-cooled water (Claim 39) or by spraying with liquid coolant (Claim 40) such as water (Claim 41) or by means of gas (Claim 42) or by means of cooled gas (Claim 43). Heyes et al. does not teach or suggest any of applicants' claims. Takano et al. does not fill in the defects of Heyes et al. so as to reach applicants' claimed invention.

The Office Action stated: that Takano et al. teaches that the shock-like cooling can be carried out by *any* known means provided that cooling conditions are satisfied (see column 6, lines 24 to 27); that water spray and partial looping of metal foil containing laminate over at least one cooled roller are well known means for shock-like cooling of the laminate, as evidenced by Levendusky et al. (U.S. Patent No. 5,919,517, column 3, lines 29 to 60; column 12, lines 23 and 24; column 14, lines 34 to 53); and that ice-cooled water and cooled gas are also well known conventional means for shock-like cooling laminates. Levendusky et al. is not stated to be part of this rejection. However, note that Levendusky et al.

applies extruded, continuous, molten polymer web to a heated metal strip.

Levendusky et al. also directs away from applicants' claimed invention.

The Office Action stated that it would have been obvious to one of ordinary skill in the art at the time the invention was made to have used any conventional cooling means including ice-cooled water, cooled gas, water spray, or partial looping over at least one cooled roller of a metal foil containing laminate shock-like cooling of a coated metal foil of Heyes et al. in view of Takano et al. since Takano et al. teaches that the shock-like cooling can be carried out by any known means provided that cooling conditions are satisfied. Applicants traverse this statement. One ordinarily skilled in the art would not have any motivation to insert Takano et al. into Heyes et al.

The Office Action stated that, as to Claims 37 and 38, Heyes et al. further teaches that shock-like cooling is carried out by passing through water (see Fig. 1). The result still would not be applicants' claimed invention.

The Office Action stated that, as to Claims 44 and 45, Heyes et al. further teaches that an adhesion promoting agent is a copolymer of propylene such as maleic anhydride (MAH) graft modified PP (see column Table I, example H). The result still would not be applicants' claimed invention.

The Office Action stated that, as to Claim 46, Claim 46 is not addressed as further limiting non-selected member of Markush group. Such claim is patentable.

The Office Action stated: that, as to Claim 47, Heyes et al. teaches that it is *preferred* to laminate coextruded hot PP-based layers to a heated aluminum

foil (see column 3, lines 40 to 45), and then re-heat the coated aluminum foil; and that Heyes et al. fails to teach that aluminum foil is at room temperature when the aluminum foil is combined with coextrudated PP-based layers. Applicants traverse this statement as being an incorrect statement of the disclosure of Heyes et al. What Heyes et al. actually discloses is (i) a generic process and (ii) two preferred subgeneric/species processes. The two preferred subgeneric/species processes are each multi-stepped with one step thereof using a metal sheet preheated to above the melting point of the polyester film (i) or preheated to between above the softening point of the polyester inner layer and below the melting point of the outer layer (ii). Column 3, lines 40 to 45, of Heyes et al. does not mention PP - it only recites polyester. No where does Heyes et al. say it is preferred to laminate coextrudaed hot PP-based layers to a heated aluminum foil (this step is one step in a preferred multi-step process). Furthermore, Heyes et al. directs away from the use of PP because it provides poor results, etc.

Also, nowhere does Heyes et al. disclose the use of hot coextruded PP-based layers. Figures 1 and 2 show rolls of polyester film being used - there is no indication that they are other than at room temperature.

The Office Action stated that Takano et al. teaches that PP-based layers on metal foil should not be allowed to cool to 100°C or lower since MAH modified PP crystallizes rapidly from about 100°C (see column 5, lines 42 to 52) to provide the coated metal foil with superior corrosion resistance and peeling resistance (see column 1, lines 57 to 62; column 2, lines 9 to 11). This information is of no

importance concerning the obviousness issue. Note that Takano et al. requires a steel sheet preheated between 100°C and 160°C. Claim 47 requires the aluminum sheet to be at room temperature. Takano et al. does not disclose the use of rollers for lamination.

The Office Action stated that common sense dictates that a temperature drop to 100°C or lower can be prevented in the process of Heyes et al. even when hot coextruded PP-based layers are laminated to an aluminum foil at room temperature if the aluminum foil enters a heater 10 shortly or right after laminating. Applicants traverse this statement for several reasons. Takano et al. states: "The temperature of the \*\*\* combination \*\*\* becomes nearly equal to the preheating temperature of the steel sheet immediately after the lamination \*\*\*."

[Column 5, lines 36 to 40] So much for common sense, which has no basis for use as such in Section 103(a) or the Graham decision requirements. Common sense is often faulty in its results, conclusions, etc. Section 103(a) requires facts. Takano et al. does not use rollers so it is not relevant to Heyes et al.

Applicants achieve laminate in a very short time, that is, almost instantaneously in the very short distance of contact point/region between the two rollers. The melted coextrudate adheres to the aluminum foil, with apparently improved adherence, almost instantaneously, with its outer surface cooling below the crystallite melt point.

The Office Action stated that it would have been obvious to one of ordinary skill in the art at the time the invention was made to have used an aluminum foil at room temperature for laminating with hot coextruded PP-based

layers as long as the temperature of the coated aluminum foil is not allowed to drop to 100°C or lower with the expectation of preventing crystallization of PP-based layers thereby providing the coated aluminum foil with the desired superior corrosion resistance and peeling resistance, as taught by Takano et al.

Applicants traverse this statement as being mere speculation based on forbidden hindsight. The bases relied on for this statement are in error.

The Office Action stated: that, as to Claims 53 to 54, Heyes et al. further teaches that coated aluminum foil made by a process comprising shock-like cooling of heated coated aluminum foil can be used for making cans (see column 1, lines 14 to 19); and that, however, Heyes et al. fails to teach that plastic coated metal cans may be used for packaging moist food including animal feed. Heyes et al. teaches one ordinarily skilled in the art not to use PP coatings so Takano et al. is not relevant.

The Office Action stated that Takano et al. teaches that PP coated aluminum foil made by a process comprising shock-like cooling of heated coated aluminum foil provides the coated aluminum foil with superior corrosion resistance and peeling resistance (see column 1, lines 57 to 62; column 2, lines 9 to 11). The Examiner has no basis in the record for combining Heyes et al. and Takano et al. No motivation to do such is present.

The Office Action stated that it would have been obvious to one of ordinary skill in the art at the time the invention was made to have used PP coated aluminum cans made by the process of Heyes et al. for packaging moist food including most animal feed since Takano et al. teaches that PP coated

0

aluminum foil made by a process comprising shock-like cooling of heated coated aluminum foil provides coated aluminum foil with superior corrosion resistance and peeling resistance. Applicants traverse this statement for the above reasons.

This rejection should be withdrawn.

The amendment filed on November 26, 2002 is objected to under 35 U.S.C. 132 on the ground that it introduces new matter into the disclosure. Applicants traverse this objection.

The Office Action stated: that 35 U.S.C. 132 states that no amendment shall introduce new matter into the disclosure of the invention; and that the added material which is not supported by the original disclosure is as follows: "The temperature of the aluminum foil, with which the coextruded plastic and adhesion-promotion agent is being combined, is such that the temperature at the surface of the plastic coating and the adhesion-promotion agent lies *below* the crystallite melt point (Tk) of the plastic." Applicant disagree that such quoted material is new matter for the reasons given above and below.

Referring to Figure 1, the distance between the outer end of the nozzle of extruder 12 and the nip region of rollers 20, 22 is small. The result of this short distance is that the reduction in temperature of melted coextrudate 14, 16 is minimal. The melted coextrudate 14, 16 and the aluminum foil met going into the nip region and are in the nip region for an instance. However, the temperature of the melted coextrudate essentially instantaneously drops to the extent that the temperature of the outer surface of the extrudate is less than the crystallite melt

point of the polyolefin 14. This is so because, if the outer surface of the olefin was at or above such melt temperature, the pressure from the two rollers would squish and disrupt or force away at least the outer portion of the coextrudate. The temperature of the aluminum foil, before and after contact with the melted coextrudate, is below the crystallite melt point of polyolefin 14. The language objected to by the Examiner is supported by the disclosure, scientific/technical principles, and the knowledge of one skilled in the art as to what would happen in the first step of applicants' claimed process as a result of the apparatus and its arrangement shown in Figure 1, for example.

The Office Action stated that applicants are required to cancel the new matter in the reply of this Office Action. Applicants traverse this requirement because it has been shown that new matter is now involved.

This objection should be withdrawn.

Claims 28 to 54 have been rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventors, at the time the application was filed, had possession of the claimed invention. Applicants traverse this rejection.

The Office Action stated that the recitation "the temperature at the surface of the plastic coating (14) and the adhesion-promotion agent (16) lies **below** the crystallite melt point (Tk) of the plastic" in independent Claim 28 is new matter since it was not described in the specification as filed. Applicant disagree with this statement and have shown above that the claim recitation is necessarily

described and supported in the specification, and by inherency, and by the knowledge of one skilled in the art.

This rejection should be withdrawn.

Claims 28 to 54 have been rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to participate point out and distinctly claim the subject matter which applicants regard as the invention. Applicants traverse this rejection.

The Office Action stated: that Claim 28, lines 6 to 7, 11 and 12, the recitation "the temperature at the surface of the plastic coating (14) and the adhesion-promotion agent (16) lies" renders the claim indefinite because it is not clear whether it is the outer non-coated surface of plastic (14) or the surface between the plastic (14) and agent (16); and that for examining purposes the phrase was interpreted as the outer non-coated surface of plastic (14). Claim 28 has been amended to eliminate this problem.

This rejection should be withdrawn.

Claim 28 has been objected to because of the following informality: "combine" in line 4 is suggested to change to "combining." This amendment has been made so this objection should be withdrawn.

Appl. No. 09/726,372 Reply to Office action of February 13, 2003

Reconsideration, reexamination and allowance of the claims are requested.

Respectfully submitted,

,

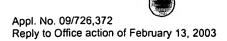
V

Virgil H. Marsh Reg. No. 23,083

Date: June 6, 2003

Fisher, Christen & Sabol 1725 K Street, N.W. Suite 1108 Washington, D.C. 20006

Tel.:(202) 659-2000 Fax:(202) 659-2015



## VERSION WITH MARKINGS TO SHOW CHANGES MADE In the Claims:

Claim 28 has been replaced with the following rewritten version of Claim 28, as amended:

28. (Once Amended) A process for production of an aluminum foil (10) coated with sealable and sterilizible plastic (14) based on polypropylene (PP) or polyethylene (PE), comprising coextruding the plastic (14) with an adhesionpromotion agent (16), to form a coextrudate, [and combine] combining the [coextruded] coextrudate of plastic (14) and adhesion-promoting agent (16) with an aluminum foil (24) between two rollers (20,22), the temperature of the coextruded-coated aluminum foil being such that the temperature at [the] outer surface of the plastic [coating] (14) of the coextrudate of the plastic (14) and the adhesion-promotion agent (16) lies below the crystallite melt point  $(T_K)$  of the plastic (14), then passing continuously the coextruded-coated aluminum foil (10), to increase the adhesion strength between the aluminum foil (24) and the plastic coating (14), through an oven (26) with temperature (To) set so that the temperature at the outer surface of the plastic coating (14) of the coextrudate of the plastic (14) and the adhesion promotion agent (16) lies above the crystallite melt point  $(T_K)$  of the plastic (14), and cooling the coextruded-coated aluminum foil (10) heat-treated in this way, after emerging from the oven (26), in a shocklike manner such that the crystalline proportion of at least in the outer surface area of the cooled plastic coating (14) and the crystal grains in the outer surface area are as small as possible.

## In the Specification:

The paragraph previously inserted on page 1, line 9, has been replaced with the following rewritten version of such paragraph on page 1, line 9, as amended:

The temperature of the aluminum foil, with which the coextrudate of the plastic and the adhesion-promotion agent is being combined with the adhesion-promotion next to the aluminum foil, is such that the temperature at the outer surface of the plastic of the coextrudate of the plastic and the adhesion-promotion agent lies below the crystallite melt point (T<sub>K</sub>) of the plastic.